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(54) Title: PROCESS TO CHIRAL BETA-AMINO ACID DERIVATIVES

(57) Abstract: The present invention relates to a process for the synthesis of chiral beta-amino acid derivatives and in particular to a process for the preparation of enantiomerically enriched beta-amino acid amide inhibitors of dipeptidyl peptidase-IV (DP-IV) which are useful for the treatment of Type 2 diabetes.

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TITLE OF THE INVENTION
PROCESS TO CHIRAL BETA-AMINO ACID DERIVATIVES

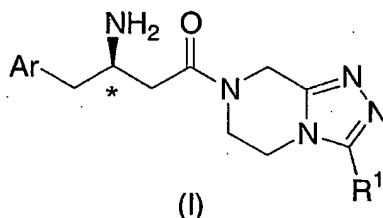
FIELD OF THE INVENTION

5 The present invention also relates to a process for the efficient preparation of enantiomerically enriched beta-amino acid derivatives which are useful in the synthesis of biologically active molecules. More particularly, the present invention relates to a process for the preparation of enantiomerically enriched beta-amino acid amide inhibitors of dipeptidyl peptidase-IV which are useful for the treatment of Type 2 diabetes.

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BACKGROUND OF THE INVENTION

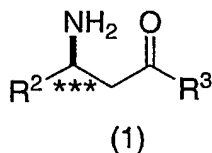
The present invention provides an efficient process for the preparation of chiral inhibitors of dipeptidyl peptidase-IV of general structural formula I,



15 or a pharmaceutically acceptable salt thereof;
having the (*R*)-configuration at the stereogenic center marked with an *.

The present invention also provides intermediates useful in the disclosed process.

The present invention also provides an efficient process for the preparation of an enantiomerically enriched beta-amino acid derivative of structural formula 1:



20

or an amine salt thereof;

having the indicated stereochemical configuration at the stereogenic center marked with an ***;
wherein

R² is C₁₋₈ alkyl, C₅₋₇ cycloalkyl, aryl, heteroaryl, aryl-C₁₋₂ alkyl, or

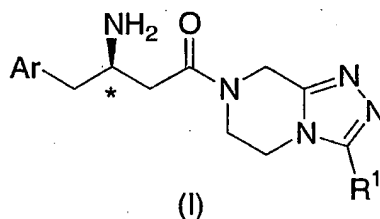
heteroaryl-C₁₋₂ alkyl, wherein aryl and heteroaryl are unsubstituted or substituted with one to three substituents independently selected from C₁₋₄ alkyl, halogen, C₁₋₄ alkoxy, and trifluoromethyl;

R³ is OR⁴, SR⁴, or NR⁴R⁵;

- 5 R⁴ and R⁵ are each independently hydrogen, C₁₋₈ alkyl, aryl, or aryl-C₁₋₂ alkyl; or R⁴ and R⁵ together with the nitrogen atom to which they are attached form a 4- to 7-membered heterocyclic ring system optionally containing an additional heteroatom selected from O, S, and NC₁₋₄ alkyl.

SUMMARY OF THE INVENTION

- 10 This invention is concerned with a process for preparing enantiomerically enriched compounds of structural formula I:



or a pharmaceutically acceptable salt thereof;

having the (*R*)-configuration at the stereogenic center marked with an *;

- 15 wherein

Ar is phenyl which is unsubstituted or substituted with one to five substituents independently selected from the group consisting of fluorine, trifluoromethyl, and trifluoromethoxy; and R¹ is hydrogen or C₁₋₄ alkyl unsubstituted or substituted with one to five fluorines.

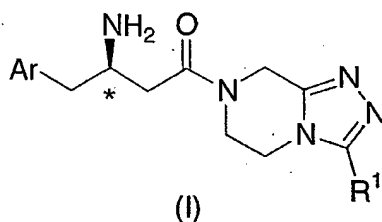
- 20 Another aspect of the present invention provides intermediate compounds of structural formulae II and IV which are useful for the preparation of compounds of structural formula I.

The products of the present process are disclosed in WO 03/004498 (published 16 January 2003) as potent inhibitors of dipeptidyl peptidase-IV which are useful for the treatment of Type 2 diabetes.

- 25 The present invention is also concerned with a process for the preparation of enantiomerically-enriched beta-amino acid derivatives of structural formula 1. The process involves the elaboration of pure *Z*-enamines from beta-ketoesters and amides using (*S*)-phenylglycine amide and their hydrogenation with very high diastereoselectivities using heterogeneous catalysis. Hydrogenolytic cleavage of the (*S*)-phenylglycine amide affords the
- 30 corresponding enantiomerically enriched beta-amino acid esters or amides.

DETAILED DESCRIPTION OF THE INVENTION

The present invention provides an efficient process for the preparation of enantiomerically enriched beta-amino acid amide derivatives of structural formula I:



5

or a pharmaceutically acceptable salt thereof;

having the (*R*)-configuration at the stereogenic center marked with an *;

wherein

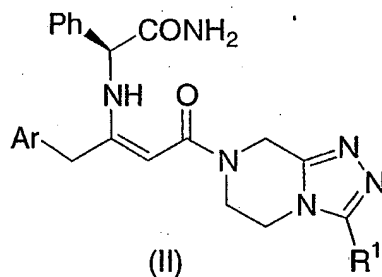
Ar is phenyl which is unsubstituted or substituted with one to five substituents independently

10 selected from the group consisting of fluorine, trifluoromethyl, and trifluoromethoxy; and

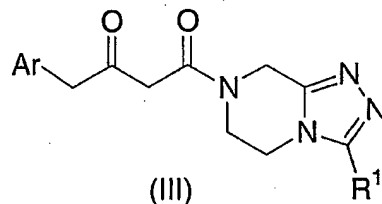
R¹ is hydrogen or C₁₋₄ alkyl unsubstituted or substituted with one to five fluorines;

comprising the steps of:

(a) producing a compound of structural formula II:

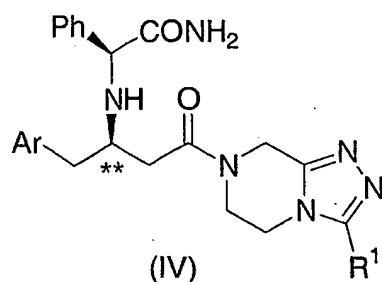


15 by treating a compound of structural formula III:

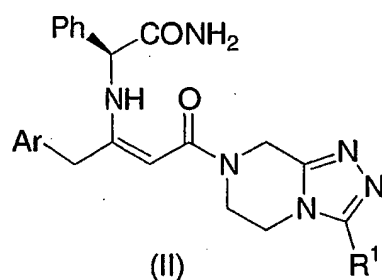


with (*S*)-phenylglycine amide in the presence of an acid in a suitable organic solvent;

(b) producing a compound of structural formula IV:

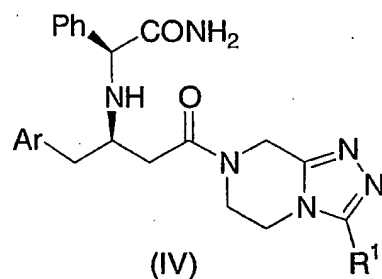


by hydrogenating a compound of structural formula II:



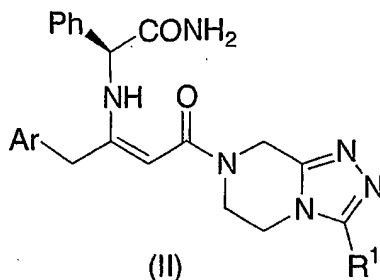
in the presence of a catalyst in a suitable organic solvent; and

5 (c) hydrogenolyzing a compound of structural formula IV:

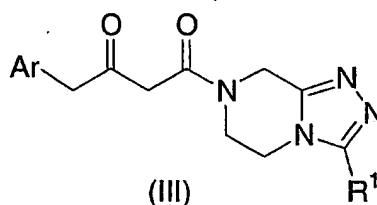


in the presence of a catalyst in a suitable organic solvent to afford a compound of structural formula I or a pharmaceutically acceptable salt thereof.

10 The first step in the process of the present invention entails the preparation of an enamine amide of structural formula II containing the (*S*)-phenylglycine amide (PGA) chiral auxiliary:



This is accomplished by treating a beta-ketoamide of formula III:



with (*S*)-phenylglycine amide in the presence of acid in a suitable organic solvent. Embodiments of acids which can be employed to generate compounds of formula II include formic acid, acetic acid, trifluoroacetic acid, methanesulfonic acid, trifluoromethanesulfonic acid, *p*-toluenesulfonic acid, and anhydrous hydrogen chloride. Suitable solvents for this step of the process include methanol, ethanol, IPA, 2,2,2-trifluoroethanol, IPAc, and mixtures thereof.

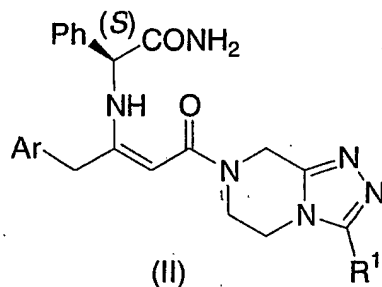
The second step in the process of the present invention concerns a diastereoselective hydrogenation of the enamine carbon-carbon double bond in the chiral substrate of formula II to afford protected chiral amines of formula IV having the (*R*)-configuration at the newly generated stereogenic center marked with an **. The hydrogenation proceeds with high diastereoselectivity (in excess of 90% de) by using platinum oxide (PtO₂) (Adam's catalyst) particularly when the catalyst is washed with acetic acid. Other catalysts that can be used include Pt/C, Pt/Al₂O₃, Pd/C, and Pd/Al₂O₃. The asymmetric hydrogenation is carried in a suitable organic solvent, such as tetrahydrofuran, a lower alkanol, for example, methanol, ethanol, IPA, and mixtures thereof, at a hydrogen gas pressure of about atmospheric pressure to about 200 psig.

The final step in the process of the present invention is removal of the (*S*)-PGA chiral auxiliary using hydrogenolytic conditions in the presence of a palladium catalyst. The hydrogenolysis can be effected with hydrogen gas or by using transfer hydrogenation conditions where hydrogen is generated *in situ*. Palladium catalysts that can be employed for the cleavage of the chiral auxiliary include Pd/C, Pd(OH)₂/C, and Pd/Al₂O₃. A preferred palladium catalyst is 20% Pd(OH)₂/C. Transfer hydrogenation reagents as sources of hydrogen gas include

cyclohexene, cyclohexadiene, formic acid, ammonium formate, tetramethylammonium formate, sodium formate, potassium formate, and isopropyl alcohol. The hydrogenolysis reaction is performed in a suitable organic solvent or aqueous organic solvent, such as THF, methanol, ethanol, IPA, 2,2,2-trifluoroethanol, IPAc, and mixtures thereof. The organic solvent may be admixed with acetic acid.

Finally, the diastereoselective hydrogenation and debenzylation can be performed in one pot by adding the palladium catalyst after the enamine hydrogenation with platinum oxide or compatible catalyst described above.

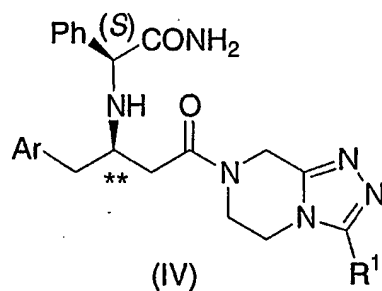
Another aspect of the process of the present invention comprises the following novel compounds of structural formula II which are intermediates in the preparation of the compounds of structural formula I:



wherein

Ar is phenyl which is unsubstituted or substituted with one to five substituents independently selected from the group consisting of fluorine, trifluoromethyl, and trifluoromethoxy; and R¹ is hydrogen or C₁₋₄ alkyl unsubstituted or substituted with one to five fluorines. In one embodiment of the novel intermediates of structural formula II, Ar is 2,5-difluorophenyl or 2,4,5-trifluorophenyl and R¹ is trifluoromethyl.

Yet a further aspect of the process of this invention comprises the following diastereomerically enriched compounds of structural formula IV which are intermediates in the preparation of the compounds of structural formula I:

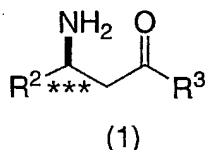


having the (*R*)-configuration at the stereogenic center marked with an **;
wherein

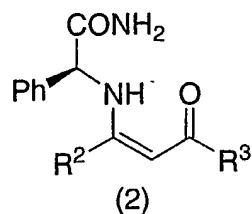
- Ar is phenyl which is unsubstituted or substituted with one to five substituents independently selected from the group consisting of fluorine, trifluoromethyl, and trifluoromethoxy; and
5 R¹ is hydrogen or C₁₋₄ alkyl unsubstituted or substituted with one to five fluorines. In one embodiment of the novel intermediates of structural formula IV, Ar is 2,5-difluorophenyl or 2,4,5-trifluorophenyl and R¹ is trifluoromethyl.

- In one embodiment of this aspect of the present invention, the *S,R*-diastereomer of structural formula IV is present in a diastereomeric excess of at least 90% over the *S,S*-diastereomer. In a class of this embodiment the *S,R*-diastereomer of structural formula IV is
10 present in a diastereomeric excess of at least 95% over the *S,S*-diastereomer.

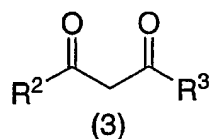
Another aspect of the present invention provides an efficient process for the preparation of enantiomerically enriched beta-amino acid derivatives of structural formula 1:



- 15 or an amine salt thereof;
having the indicated stereochemical configuration at the stereogenic center marked with an ***;
wherein
R² is C₁₋₈ alkyl, C₅₋₇ cycloalkyl, aryl, heteroaryl, aryl-C₁₋₂ alkyl, or heteroaryl-C₁₋₂ alkyl, wherein aryl and heteroaryl are unsubstituted or substituted with one to
20 three substituents independently selected from C₁₋₄ alkyl, halogen, C₁₋₄ alkoxy, and trifluoromethyl;
R³ is OR⁴, SR⁴, or NR⁴R⁵;
R⁴ and R⁵ are each independently hydrogen, C₁₋₈ alkyl, aryl, or aryl-C₁₋₂ alkyl; or R⁴ and R⁵ together with the nitrogen atom to which they are attached form a 4- to 7-membered heterocyclic
25 ring system optionally containing an additional heteroatom selected from O, S, and NC₁₋₄ alkyl;
comprising the steps of:
(a) producing a compound of structural formula 2:

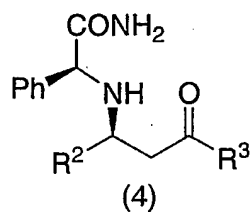


by treating a compound of structural formula 3:

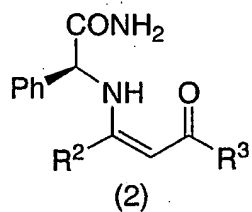


with (*S*)-phenylglycine amide in the presence of an acid in a suitable organic solvent;

5 (b) producing a compound of structural formula 4:

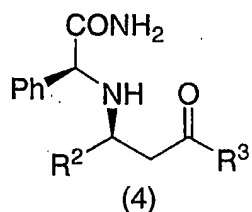


by hydrogenating a compound of structural formula 2:



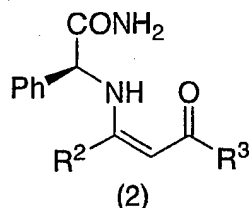
in the presence of a catalyst in a suitable organic solvent; and

10 (c) hydrogenolyzing a compound of structural formula 4:

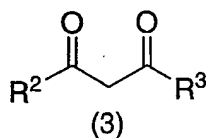


in the presence of a catalyst in a suitable organic solvent to afford a compound of structural formula 1 or a salt thereof.

The first step in the process of the present invention entails the preparation of an enamine ester or amide of structural formula 2 containing the (*S*)-phenylglycine amide (PGA) chiral auxiliary:



- 5 This is accomplished by treating a beta-ketoester or amide of formula 3:



- with (*S*)-phenylglycine amide in the presence of acid in a suitable organic solvent. Embodiments of acids which can be employed to generate compounds of formula 2 include formic acid, acetic acid, trifluoroacetic acid, methanesulfonic acid, trifluoromethanesulfonic acid, *p*-toluenesulfonic acid, and anhydrous hydrogen chloride. A preferred acid is acetic acid which can be used in catalytic amounts. Suitable solvents for this step of the process include methanol, ethanol, isopropanol (IPA), 2,2,2-trifluoroethanol, isopropyl acetate, and mixtures thereof. Preferred solvents are methanol and isopropanol.

- The second step in the process to compounds of formula 1 concerns a diastereoselective hydrogenation of the enamine carbon-carbon double bond in the chiral substrate of formula 2 to afford protected chiral amines of formula 4 having the indicated stereochemical configuration at the newly generated stereogenic center marked with an ***. The hydrogenation proceeds with high diastereoselectivity (in excess of 90% de) by using platinum oxide (PtO₂) (Adam's catalyst) particularly when the catalyst is washed with acetic acid. The asymmetric hydrogenation is carried in a suitable organic solvent, such as tetrahydrofuran, a lower alkanol, for example, methanol, ethanol, IPA, and mixtures thereof, at a hydrogen gas pressure of about atmospheric pressure to about 200 psig. A preferred organic solvent is tetrahydrofuran.

- The final step in the process of the present invention is removal of the (*S*)-PGA chiral auxiliary using hydrogenolytic conditions in the presence of a palladium catalyst. The hydrogenolysis can be effected with hydrogen gas or by using transfer hydrogenation conditions where hydrogen is generated *in situ*. Palladium catalysts that can be employed for the cleavage

of the chiral auxiliary include Pd/C, Pd(OH)₂/C, and Pd/Al₂O₃. A preferred palladium catalyst is 20% Pd(OH)₂/C. Transfer hydrogenation reagents as sources of hydrogen gas include cyclohexene, cyclohexadiene, formic acid, ammonium formate, tetramethylammonium formate, sodium formate, potassium formate, and isopropyl alcohol. The hydrogenolysis reaction is performed in a suitable organic solvent or aqueous organic solvent, such as THF, methanol, ethanol, IPA, 2,2,2-trifluoroethanol, IPAc, and mixtures thereof. The organic solvent may be admixed with acetic acid.

Finally, the diastereoselective hydrogenation and debenzylation can be performed in one pot by adding the palladium catalyst after the enamine hydrogenation with platinum oxide or compatible catalyst described above.

In one embodiment of this aspect of the present invention, R³ is 2,4,5-trifluorobenzyl and R² is C₁₋₄ alkoxy. In a class of this embodiment, R² is methoxy.

Throughout the instant application, the following terms have the indicated meanings:

The term "% enantiomeric excess" (abbreviated "ee") shall mean the % major enantiomer less the % minor enantiomer. Thus, an 80% enantiomeric excess corresponds to formation of 90% of one enantiomer and 10% of the other. The term "enantiomeric excess" is synonymous with the term "optical purity."

The term "enantiomerically enriched" shall mean that a compound of structural formula I is obtained by the process of the present invention with an enantiomeric excess of the desired (*R*)-enantiomer greater than 70% over the (*S*)-enantiomer. In one embodiment a compound of formula I having the (*R*)-configuration is obtained with an ee greater than 80%. In a class of this embodiment the (*R*)-enantiomer is obtained with an ee greater than 90%. In a subclass of this class the (*R*)-enantiomer is obtained with an ee greater than 95%.

The term "% diastereomeric excess" (abbreviated "de") shall mean the % major diastereomer less the % minor diastereomer. Thus, an 80% diastereomeric excess corresponds to formation of 90% of one diastereomer and 10% of the other.

The term "enantioselective" shall mean a reaction in which one enantiomer is produced (or destroyed) more rapidly than the other, resulting in the predominance of the favored enantiomer in the mixture of products.

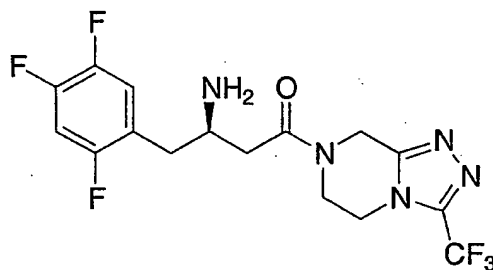
The term "diastereoselective" shall mean a reaction in which one diastereomer is produced (or destroyed) more rapidly than the other, resulting in the predominance of the favored diastereomer in the mixture of products.

Representative experimental procedures utilizing the novel process are detailed below. The following Examples are provided for the purpose of illustration only, but doing so is

not intended to limit the process of the present invention to the specific conditions for making these particular compounds.

Abbreviations: AcOH is acetic acid; IPA is isopropyl alcohol; IPAc is isopropyl acetate; THF is tetrahydrofuran; TFE is 2,2,2-trifluoroethanol; DCM is dichloromethane; DMSO is dimethylsulfoxide; MeOH is methanol.

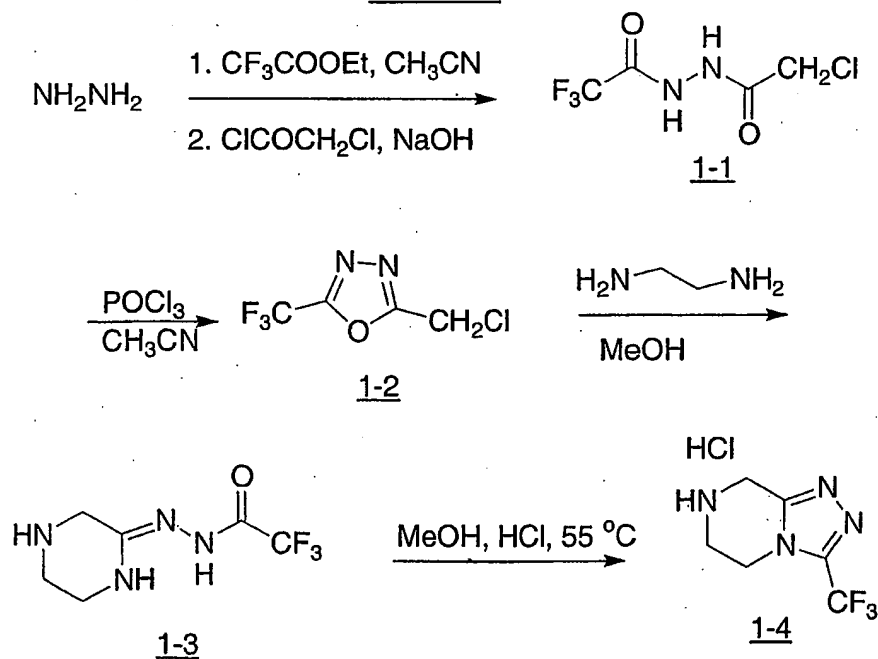
EXAMPLE 1



10 (2R)-4-oxo-4-[3-(trifluoromethyl)-5,6-dihydro[1,2,4]triazolo[4,3-a]pyrazin-7(8H)-yl]-1-(2,4,5-trifluorophenyl)butan-2-amine (2-6)

I. Preparation of 3-(trifluoromethyl)-5,6,7,8-tetrahydro[1,2,4]triazolo[4,3-a]pyrazine, hydrochloride salt (1-4)

Scheme 1



15

Step A: Preparation of bishydrazide (1-1)

Hydrazine (20.1 g, 35 wt% in water, 0.22 mol) was mixed with 310 mL of acetonitrile. 31.5 g of ethyl trifluoroacetate (0.22 mol) was added over 60 min. The internal temperature was increased to 25 °C from 14 °C. The resulting solution was aged at 22 - 25 °C for 60 min. The solution was cooled to 7 °C. 17.9 g of 50 wt% aqueous NaOH (0.22 mol) and 25.3 g of chloroacetyl chloride (0.22 mol) were added simultaneously over 130 min at a temperature below 16 °C. When the reaction was complete, the mixture was vacuum distilled to remove water and ethanol at 27 ~ 30 °C and under 26 ~ 27 in Hg vacuum. During the distillation, 720 mL of acetonitrile was added slowly to maintain constant volume (approximately 500 mL). The slurry was filtered to remove sodium chloride. The cake was rinsed with about 100 mL of acetonitrile. Removal of the solvent afforded bis-hydrazide 1-1 (43.2 g, 96.5% yield, 94.4 area% pure by HPLC assay).

¹H-NMR (400 MHz, DMSO-*d*₆): δ 4.2 (s, 2H), 10.7 (s, 1H), and 11.6 (s, 1H) ppm.

¹³C-NMR (100 MHz, DMSO-*d*₆): δ 41.0, 116.1 (q, J = 362 Hz), 155.8 (q, J = 50 Hz), and 165.4 ppm.

Step B: Preparation of 5-(trifluoromethyl)-2-(chloromethyl)-1,3,4-oxadiazole (1-2)

Bishydrazide 1-1 from Step A (43.2 g, 0.21 mol) in ACN (82 mL) was cooled to 5 °C. Phosphorus oxychloride (32.2 g, 0.21 mol) was added, maintaining the temperature below 10 °C. The mixture was heated to 80 °C and aged at this temperature for 24 h until HPLC showed less than 2 area% of 1-1. In a separate vessel, 260 mL of IPAc and 250 mL of water were mixed and cooled to 0 °C. The reaction slurry was charged to the quench keeping the internal temperature below 10 °C. After the addition, the mixture was agitated vigorously for 30 min, the temperature was increased to room temperature and the aqueous layer was cut. The organic layer was then washed with 215 mL of water, 215 mL of 5 wt% aqueous sodium bicarbonate and finally 215 mL of 20 wt% aqueous brine solution. HPLC assay yield after work up was 86-92%. Volatiles were removed by distillation at 75-80 mm Hg, 55 °C to afford an oil which could be used directly in Step C without further purification. Otherwise the product can be purified by distillation to afford 1-2 in 70-80% yield.

¹H-NMR (400 MHz, CDCl₃): δ 4.8 (s, 2H) ppm.

¹³C-NMR (100 MHz, CDCl₃): δ 32.1, 115.8 (q, J = 337 Hz), 156.2 (q, J = 50 Hz), and 164.4 ppm.

Step C: Preparation of N-[(2Z)-piperazin-2-ylidene]trifluoroacetohydrazide (1-3)

To a solution of ethylenediamine (33.1 g, 0.55 mol) in methanol (150 mL) cooled at -20 °C was added distilled oxadiazole 1-2 from Step B (29.8 g, 0.16 mol) while keeping the

internal temperature at -20°C . After the addition was complete, the resulting slurry was aged at -20°C for 1 h. Ethanol (225 mL) was then charged and the slurry slowly warmed to -5°C .

After 60 min at -5°C , the slurry was filtered and washed with ethanol (60 mL) at -5°C .

Amidine 1-3 was obtained as a white solid in 72% yield (24.4 g, 99.5 area wt% pure by HPLC).

5 ^1H -NMR (400 MHz, $\text{DMSO}-d_6$): δ 2.9 (t, 2H), 3.2 (t, 2H), 3.6 (s, 2H), and 8.3 (b, 1H) ppm.

^{13}C -NMR (100 MHz, $\text{DMSO}-d_6$): δ 40.8, 42.0, 43.3, 119.3 (q, $J = 350$ Hz), 154.2, and 156.2 (q,

$J = 38$ Hz) ppm.

Step D: Preparation of 3-(trifluoromethyl)-5,6,7,8-tetrahydro[1,2,4]triazolo[4,3-
a]pyrazine, hydrochloride salt (1-4)

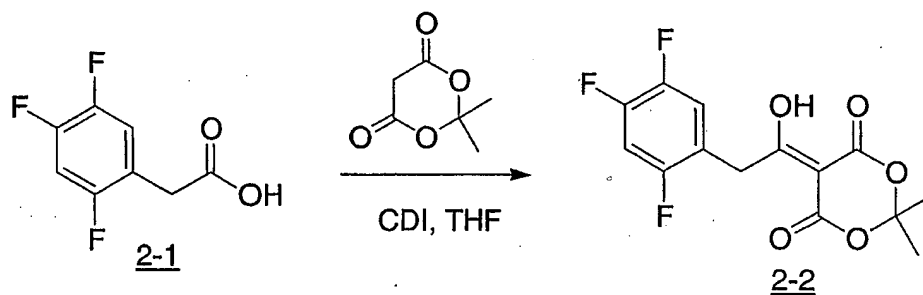
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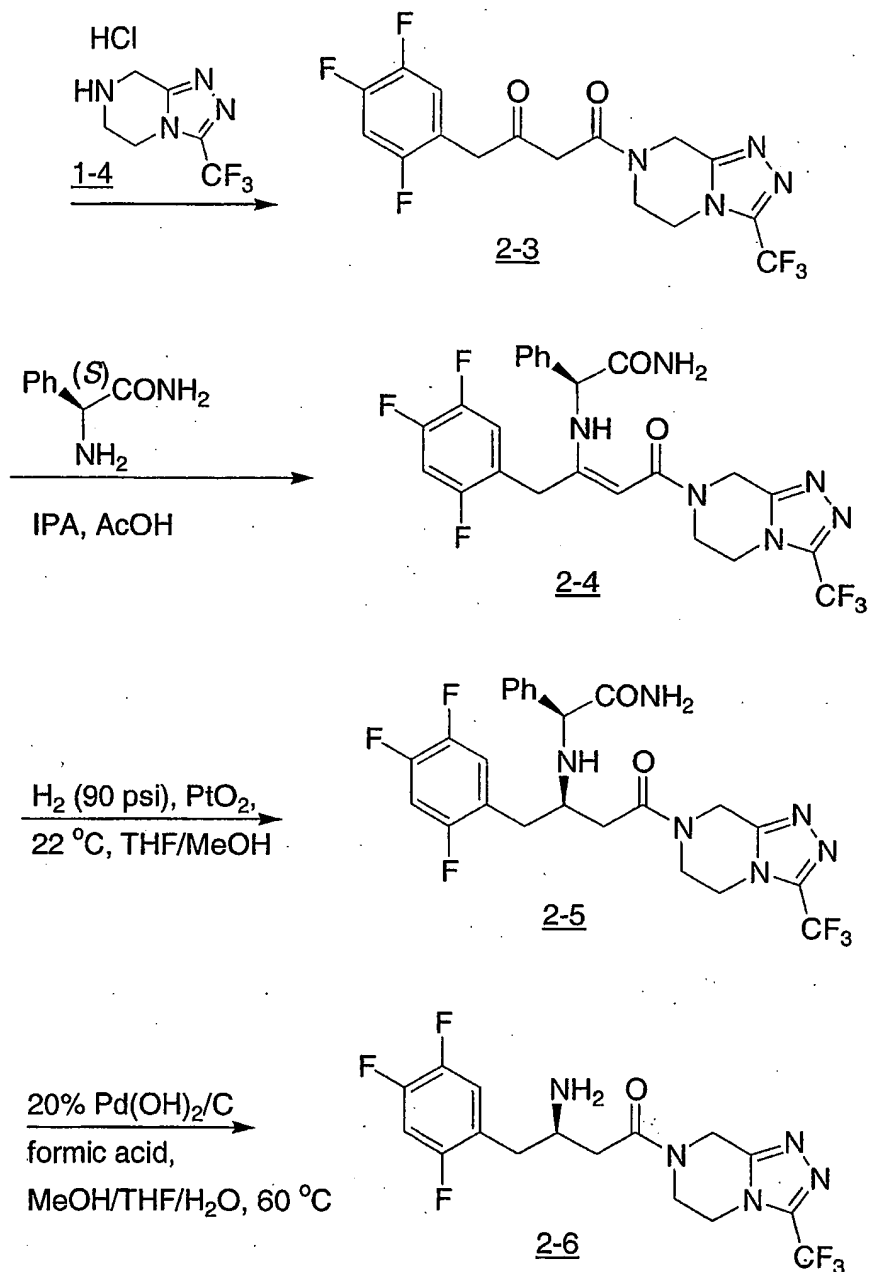
A suspension of amidine 1-3 (27.3 g, 0.13 mol) in 110 mL of methanol was warmed to 55°C . 37% Hydrochloric acid (11.2 mL, 0.14 mol) was added over 15 min at this temperature. During the addition, all solids dissolved resulting in a clear solution. The reaction was aged for 30 min. The solution was cooled down to 20°C and aged at this temperature until
15 a seed bed formed (10 min to 1 h). 300 mL of MTBE was charged at 20°C over 1 h. The resulting slurry was cooled to 2°C , aged for 30 min and filtered. Solids were washed with 50 mL of ethanol:MTBE (1:3) and dried under vacuum at 45°C . Yield of triazole 1-4 was 26.7 g (99.5 area wt% pure by HPLC).

^1H -NMR (400 MHz, $\text{DMSO}-d_6$): δ 3.6 (t, 2H), 4.4 (t, 2H), 4.6 (s, 2H), and 10.6 (b, 2H) ppm;

20 ^{13}C -NMR (100 MHz, $\text{DMSO}-d_6$): δ : 39.4, 39.6, 41.0, 118.6 (q, $J = 325$ Hz), 142.9 (q, $J = 50$ Hz), and 148.8 ppm.

Scheme 2





5 Step A: Preparation of 5-[1-hydroxy-2-(2,4,5-trifluorophenyl)ethylidene]-2,2-dimethyl-1,3-dioxane-4,6-dione (2-2)

2,4,5-Trifluorophenylacetic acid (2-1) (11.4 g, 60 mmol) was dissolved in THF (60 mL) and 1,1'-carbonyldiimidazole (10.7 g, 66 mmol) was added over 5 min. The mixture was warmed to 51°C , Meldrum's acid (9.51 g, 66 mmol) was added, and the mixture was aged
 10 for 3 h. The reaction mixture was diluted with IPAc (60 mL) and water (60 mL), and the pH

was adjusted to 2.4 with concentrated hydrochloric acid (11.5 mL). The aqueous layer was separated, and the organic layer was washed at 36 °C with 0.1 N HCl (60 mL). The organic layer was concentrated, flushed with IPAc, and the residue was slurried in 2:1 heptane/IPAc (70 mL). The mixture was cooled over an ice-bath, then filtered, rinsing the solids with 2:1
5 heptane/IPAc. After drying, the Meldrum's acid adduct 2-2 was obtained as a solid (15.1 g).

Step B: Preparation of 4-oxo-4-[3-(trifluoromethyl)-5,6-dihydro[1,2,4]triazolo[4,3-
a]pyrazin-7(8H)-yl]-1-(2,4,5-trifluorophenyl)butan-2-one (2-3)

The Meldrum's acid adduct 2-2 from Step A (22.1 g, 70 mmol) and the triazole
10 hydrochloride 1-4 (16.0 g, 70 mmol) were slurried in IPAc (220 mL) and *N,N*-
diisopropylethylamine (12.8 mL) was added. After aging for 3.5 h at 85 °C, water (175 mL) was
added and the mixture was transferred to a separatory funnel with a 40-mL rinse with IPAc. The
aqueous layer was separated and the organic layer was washed with water (100 mL). The
organic layer was partially concentrated under diminished pressure to give a solution of
15 ketoamide 2-3 (65 g) in IPAc. *n*-Heptane (30 mL) was added at room temperature, followed by
seed crystals of ketoamide. More heptane (20 mL) was added dropwise, and the mixture was
stirred overnight. Then more heptane (50 mL) was added slowly and after aging for 2 h, the
solids were filtered and washed with 2.2:1 heptane/IPAc (30 mL). After drying, the ketoamide
2-3 was obtained in 92% yield (26.3 g).

Step C: Preparation of (2S)-2-((1Z)-3-oxo-1-(2,4,5-trifluorobenzyl)-3-[3-
(trifluoromethyl)-5,6-dihydro[1,2,4]triazolo[4,3-a]pyrazin-7(8H)-yl]prop-1-
enyl)amino)-2-phenylethanamide (2-4)

Ketoamide 2-3 (98.4 wt%, 711 g, 1.72 mol) and (*S*)-phenylglycine amide (98.4
25 wt%, 276 g, 1.81 mol) were added to 2.8 L IPA, warmed to 40 °C and AcOH (49 mL) was
added. The temperature rose initially to 48 °C after 15 min and came down to 40 °C in 1 h.
After aging for 5 h, 0.4% seed was added and the mixture was aged 1 h to afford a slurry. This
mixture was distilled at constant volume (39 °C, 98 Torr) with 2.0 L IPA flush over 2.5 h. The
mixture was aged for 1 h and 2.8 L heptane was added over 3 h. The mixture was aged 3.5 h at
30 40 °C and cooled down slowly to room temperature over 5.5 h. After 21 h total reaction time,
the slurry was rapidly filtered and rinsed with 800 mL 1:1 IPA/heptane. The solid was dried for
24 h under N₂ to afford 972 g of PGA-enamine 2-4 (86.6 wt%, 98.7 area% purity, 91% yield) as
an IPA solvate.

Step D: Preparation of (2S)-2-(((1R)-3-oxo-1-(2,4,5-trifluorobenzyl)-3-[3-(trifluoromethyl)-5,6-dihydro[1,2,4]triazolo[4,3-a]pyrazin-7(8H)-yl]propyl}amino)-2-phenylethanamide (2-5)

Intermediate 2-4 (20.0 g) and PtO₂ (0.500 g) were suspended in 80 mL THF and 20 mL MeOH (22 °C) in a 300 mL stirred autoclave. The stirred solution was cooled to 15 °C and placed under hydrogen pressure (90 psig). After 30 min, the solution was warmed to 22 °C over 30 min and aged for 26 h. The autoclave was vented to atmospheric pressure and the reaction solution was filtered through Solka floc. The filter cake was rinsed with methanol (2 x 20 mL). The filtrate and washings were combined and carried on directly to the final debenzylation Step E. Compound 2-5 was obtained in 90% assay yield (about 17.8 g) and 96.4% diastereomeric excess (de).

The PtO₂ catalyst was prepared as follows: PtO₂ (36.7 g) was suspended in acetic acid (130 mL). The stirred slurry was aged at room temperature for 2 h and then filtered. The filter cake was rinsed with acetic acid (4 x 25 mL) and then dried *in vacuo* at 50 °C for 24 h. The PtO₂ was isolated with 97% recovery (35.4 g).

Step E: Preparation of (2R)-4-oxo-4-[3-(trifluoromethyl)-5,6-dihydro[1,2,4]triazolo[4,3-a]pyrazin-7(8H)-yl]-1-(2,4,5-trifluorophenyl)butan-2-amine (2-6)

The crude product from Step D (41.5 g, 76.3 mmol) and 20% Pd(OH)₂/C (12.4 g, 30 wt%) was slurried in 1:1 THF/MeOH (124 mL) and water (41.5 mL) and formic acid (41.5 mL) were added. The reaction mixture was heated at 60 °C for 3 h. After cooling, the reaction was treated with Solka floc (10 g) and filtered through Solka floc (20 g), rinsing with MeOH (200 mL). The filtrate contained 29.4 g (94.5% assay yield) of 2-6 with an optical purity of 97% ee.

Alternative Step E:

The crude product from Step D (82.5 g, 153 mmol) was dissolved in 1:1 THF/MeOH (200 mL) and finely pulverized 20% Pd(OH)₂/C (24.8 g, 30 wt%) was added. The mixture was added to a stainless-steel autoclave. THF/MeOH (130 mL) was used to rinse the mixture into the vessel, then acetic acid (21.9 mL) and water (82.5 mL) were added. The mixture was heated at 50 °C for 10 h at 40 psig hydrogen. After cooling to room temperature and venting to ambient pressure, the autoclave was emptied with MeOH rinse (2 x 100 mL). The batch was treated with Solka floc (11 g) and filtered through Solka floc (42 g). The filter pad was washed with MeOH (200 mL). The filtrate contained 57.9 g of 2-6 with an optical purity of 96% ee.

The optical purity of 2-6 was further enhanced in the following manner. The solution from the hydrogenation reaction (18 g in 180 mL solvent) was concentrated and switched to methyl *t*-butyl ether (MTBE) (45 mL). Into this solution was added aqueous H₃PO₄ solution (0.5 M, 95 mL). After separation of the layers, 3N NaOH (35 mL) was added to the water layer, which was then extracted with MTBE (180 mL + 100 mL). The MTBE solution was concentrated and solvent switched to hot toluene (180 mL, about 75 °C). The hot toluene solution was then allowed to cool to 0 °C slowly (5 – 10 h). The crystals were isolated by filtration (13 g, yield 72%, 98 – 99% ee); m.p. 114.1 – 115.7 °C.

¹H NMR (300 MHz, CD₃CN): δ 7.26 (m), 7.08 (m), 4.90 (s), 4.89 (s), 4.14 (m), 3.95 (m), 3.40 (m), 2.68 (m), 2.49 (m), 1.40 (bs).

Compound 2-6 exists as amide bond rotamers. Unless indicated, the major and minor rotamers are grouped together since the carbon-13 signals are not well resolved:

¹³C NMR (CD₃CN): δ 171.8, 157.4 (ddd, *J*_{CF} = 242.4, 9.2, 2.5 Hz), 152.2 (major), 151.8 (minor), 149.3 (ddd; *J*_{CF} = 246.7, 14.2, 12.9 Hz), 147.4 (ddd, *J*_{CF} = 241.2, 12.3, 3.7 Hz), 144.2 (q, *J*_{CF} = 38.8 Hz), 124.6 (ddd, *J*_{CF} = 18.5, 5.9, 4.0 Hz), 120.4 (dd, *J*_{CF} = 19.1, 6.2 Hz), 119.8 (q, *J*_{CF} = 268.9 Hz), 106.2 (dd, *J*_{CF} = 29.5, 20.9 Hz), 50.1, 44.8, 44.3 (minor), 43.2 (minor), 42.4, 41.6 (minor), 41.4, 39.6, 38.5 (minor), 36.9.

The following high-performance liquid chromatographic (HPLC) conditions were used to determine percent conversion to product:

Waters Symmetry Shield RP8, 4.6 x 250 mm, 5 μm, 25 °C, 210 nm detection, 5 μL injection, 1 mL/min, 15 min run time, isocratic at 45% acetonitrile/55% 10 mM pH 6.8 phosphate buffer.

Retention times:

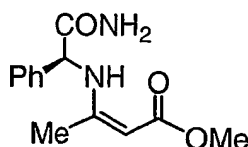
Compound <u>2-6</u> :	4.30 min
Compound <u>2-5</u> (<i>S,R</i> -diastereomer):	7.73 min
Compound <u>2-5</u> (<i>S,S</i> -diastereomer):	7.11 min

The following high-performance liquid chromatographic (HPLC) conditions were used to determine optical purity (percentage ee):

Chiralcel AD-H, 5 μm, 4.6 x 250 mm, 5 μm, 35 °C, 268 nm detection, 10 μL injection, 0.7 mL/min, 25 min run time, isocratic at 40% 0.1% diethylamine in hexane/60% 0.1% diethylamine in ethanol.

Retention times:

Compound <u>2-6</u> (<i>R</i> -enantiomer):	18.4 min
Compound <u>2-6</u> (<i>S</i> -enantiomer):	15.4 min

Examples of Z-PGA-Enamines:EXAMPLE 2Methyl (2Z)-3-[[[(1S)-2-amino-2-oxo-1-phenylethyl]amino]but-2-enoate

5

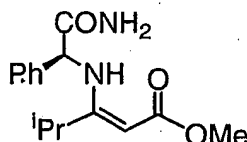
Methyl acetoacetate (2.0 g, 17.2 mmol) and (S)-PGA (2.58 g, 17.2 mmol) were dissolved in MeOH (200 mL) at 40 °C and acetic acid (0.5 mL, 8.6 mmol) was added. Solids formed after 1 h at 40 °C. The mixture was cooled to 4 °C and filtered. The solids were rinsed with cold MeOH and dried to afford the enamine (3.3 g). $[\alpha]_D^{25} = -92^\circ$ (c 0.44, DCM); mp = 159 °C.

10

$^1\text{H-NMR}$ (DMSO- d_6 , 400 MHz): δ 9.46 (d, $J = 8$ Hz, 1H), 7.70 (br, 1H), 7.37 (m, 4H), 7.28 (m, 2H), 5.19 (d, $J = 8$ Hz, 1H), 4.44 (s, 1H), 3.52 (s, 3H), 1.73 (s, 3H).

$^{13}\text{C-NMR}$ (DMSO- d_6 , 100 MHz): δ 171.2, 169.5, 160.1, 139.8, 128.7, 127.8, 126.4, 83.1, 59.0, 49.6, 19.4

15

EXAMPLE 3Methyl (2Z)-3-[[[(1S)-2-amino-2-oxo-1-phenylethyl]amino]-4-methylpent-2-enoate

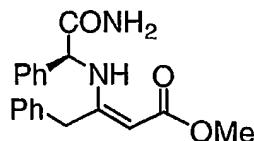
Methyl isobutyryl acetate (7.59 g, 50.0 mmol) and (S)-PGA (7.51 g, 50.0 mmol) were mixed in IPA (40 mL) at 40 °C and acetic acid (1.43 mL, 25 mmol) was added. A slurry formed quickly. The mixture was stirred overnight at 40 °C, cooled to 3 °C and was then filtered. The solids were rinsed with IPA and dried to afford the enamine as a white solid (11.6 g). $[\alpha]_D^{25} = -57^\circ$ (c 0.58, DCM); mp = 170 °C.

20

$^1\text{H-NMR}$ (DMSO- d_6 , 400 MHz): δ 9.55 (d, $J = 8$ Hz, 1H), 7.76 (s, 1H), 7.40 (m, 4H), 7.30 (m, 2H), 5.29 (d, $J = 8$ Hz, 1H), 4.48 (s, 1H), 3.54 (s, 3H), 2.44 (m, 1H), 1.08 (d, $J = 7$ Hz, 3H), 0.78 (d, $J = 7$ Hz, 3H).

25

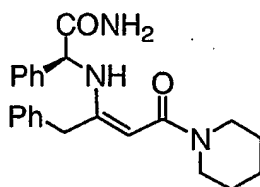
$^{13}\text{C-NMR}$ (DMSO- d_6 , 100 MHz): δ 171.5, 170.5, 170.2, 140.6, 129.0, 128.1, 126.6, 79.2, 58.5, 49.9, 29.2, 22.4, 21.1.

EXAMPLE 4Methyl (2Z)-3-[[[(1S)-2-amino-2-oxo-1-phenylethyl]amino]-4-phenylbut-2-enoate

5 Methyl 3-oxo-4-phenylbutanoate (1.76 g, 9.16 mmol) and (*S*)-PGA (1.38 g, 9.16 mmol) were mixed in isopropanol (IPA) (18 mL) and acetic acid (0.26 mL, 4.6 mmol) and stirred 18 h over a 50 °C bath. The resulting slurry was cooled and filtered. The solids were rinsed with cold MeOH and dried to afford the enamine (2.20 g).

$[\alpha_D]^{25} = -63^\circ$ (c 0.40, DCM); mp = 169 °C.

10 $^1\text{H-NMR}$ (DMSO- d_6 , 400 MHz): δ 9.46 (d, $J = 8$ Hz, 1H), 7.71 (s, 1H), 7.34 (d, 4H), 7.26 (m, 4H), 7.20 (m, 1H), 7.08 (d, 2H), 5.10 (d, $J = 8$ Hz, 1H), 4.30 (s, 1H), 3.50 (s, 3H), 3.32 (q, 2H).
 $^{13}\text{C-NMR}$ (DMSO- d_6 , 100 MHz): δ 170.8, 169.4, 161.7, 139.9, 136.3, 128.7, 128.6, 128.6, 127.8, 126.7, 126.3, 84.6, 58.6, 49.7, 37.9

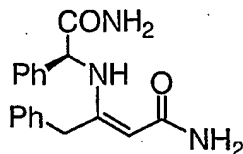
EXAMPLE 5(2S)-2-[[[(1Z)-1-benzyl-3-oxo-3-piperidin-1-ylprop-1-en-1-yl]amino]-2-phenylacetamide

20 4-Oxo-1-phenyl-4-piperidin-1-ylbutan-2-one (13.8 g, 56.0 mmol) and (*S*)-PGA (8.42 g, 56.0 mmol) were stirred in MeOH (70 mL) with acetic acid (1.6 mL, 28 mmol) 3 days at 40 °C. The mixture was concentrated and the residue was purified by flash chromatography (80-100% EtOAc/ hexane) and then was crystallized (EtOAc/ hexane) to afford the enamine (7.4 g).

$[\alpha_D]^{25} = -86^\circ$ (c 0.44, MeOH); mp = 155 °C.

25 $^1\text{H-NMR}$ (DMSO- d_6 , 400 MHz): δ 10.23 (d, $J = 9$ Hz, 1H), 7.60 (br, 1H), 7.29 (m, 9H), 7.19 (m, 1H), 7.10 (br, 1H), 4.99 (d, $J = 9$ Hz, 1H), 4.81 (s, 1H), 3.32-3.35 (m, 6H), 2.49 (m, 2H), 1.41 (m, 4H).

$^{13}\text{C-NMR}$ (DMSO- d_6 , 100 MHz): δ 171.3, 168.3, 158.2, 140.3, 137.3, 128.4, 128.3, 127.5, 126.5, 126.4, 85.1, 58.9, 38.6, 25.8, 24.3.

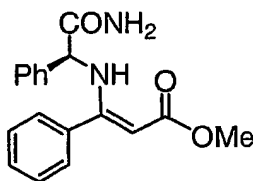
EXAMPLE 6(2Z)-3-[[[(1S)-2-amino-2-oxo-1-phenylethyl]amino]-4-phenylbut-2-enamide

3-Oxo-4-phenylbutanamide (1.85 g, 10.4 mmol) and (*S*)-PGA (1.56 g, 10.4 mmol) were stirred in MeOH (18 mL) with acetic acid (0.30 mL, 5.2 mmol) 4 days at 40 °C. The mixture was diluted with methyl tert-butyl ether (MTBE), washed with saturated sodium bicarbonate, dried (MgSO₄) and concentrated. The residue was purified by flash chromatography (50-100% EtOAc/hexane) and then crystallized (EtOAc/ hexane) to afford the enamine (0.63 g).

[α]_D²⁵ = +9° (c 0.18, DCM); mp = 188 °C.

¹H-NMR (DMSO-d₆, 400 MHz): δ 9.89 (d, *J* = 8.8 Hz, 1H), 7.61 (s, 1H), 7.33 (m, 4H), 7.28 (d, *J* = 7 Hz, 3 H), 7.21 (m, 1H), 7.15 (d, 7 Hz, 2H), 7.10 (s, 1H), 6.6 (br, 1H), 6.1 (br, 1H), 5.01 (d, *J* = 8.8 Hz, 1H), 4.37 (s, 1H), 3.27 (q, 2H).

¹³C-NMR (DMSO-d₆, 100 MHz): δ 171.7, 171.4, 157.2, 140.5, 137.2, 128.5, 128.4, 127.4, 126.4, 126.4, 89.4, 58.9, 38.1.

EXAMPLE 7Methyl (2Z)-3-[[[(1S)-2-amino-2-oxo-1-phenylethyl]amino]-3-phenylacrylate

Methyl 3-oxo-3-phenylpropanoate (20.2 g, 114 mmol) and (*S*)-PGA (17.0 g, 114 mmol) were added to MeOH (60 mL) with acetic acid (3.25 mL, 57 mmol) and stirred 4 days over a 40 °C bath. The mixture was concentrated and the residue was purified by flash chromatography (40-60% EtOAc/ hexane) to afford an oil, which was crystallized in 20% EtOAc/ hexane (250 mL) to afford after drying the enamine (19.7 g).

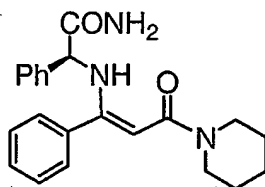
[α]_D²⁵ = -68° (c 0.49, DCM); mp = 142 °C.

¹H-NMR (DMSO-d₆, 400 MHz): δ 9.46 (d, *J* = 9 Hz, 1H), 7.65 (s, 1H), 7.39 (m, 1H), 7.33 (m, 2H), 7.24 (m, 4H), 7.11 (m, 4H), 4.92 (d, *J* = 9 Hz, 1H), 4.51 (s, 1H), 3.61 (s, 3H).

^{13}C -NMR (DMSO- d_6 , 100 MHz): δ 171.0, 169.0, 162.5, 140.0, 135.5, 129.2, 128.2, 128.2, 127.3, 127.2, 126.1, 86.4, 59.9, 49.8.

EXAMPLE 8

5 (2S)-2-[[[(1Z)-3-oxo-1-phenyl-3-piperidin-1-ylprop-1-en-1-yl]amino]-2-phenylacetamide



3-Oxo-1-phenyl-3-piperidin-1-ylpropan-1-one (10.1 g, 43.7 mmol) and (S)-PGA (6.6 g, 44 mmol) in IPA (40 mL) and acetic acid (1.9 mL, 33 mmol) were stirred 4 days at 40 °C. The mixture was concentrated and purified by flash chromatography and crystallization (EtOAc/hexane) to afford the enamine (6.9 g).

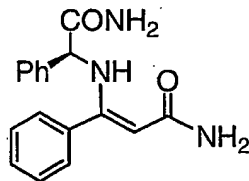
$[\alpha]_D^{25} = +86^\circ$ (c 0.26, DCM); mp = 126 °C.

^1H -NMR (DMSO- d_6 , 400 MHz): δ 10.17 (d, J = 9 Hz, 1H), 7.56 (s, 1H), 7.07-7.37 (m, 11H), 4.84 (s, 1H), 4.80 (d, J = 9 Hz, 1H), 3.41 (br, 4H), 1.56 (br, 2H), 1.33 (br, 4H).

^{13}C -NMR (DMSO- d_6 , 100 MHz): δ 171.5, 167.7, 160.0, 140.5, 137.0, 128.8, 128.2, 128.1, 127.5, 127.2, 126.4, 87.4, 60.4, 25.8, 24.3

EXAMPLE 9

(2Z)-3-[[[(1S)-2-amino-2-oxo-1-phenylethyl]amino]-3-phenylacrylamide



20 3-Oxo-3-phenylpropanamide (6.53 g, 40.0 mmol) and (S)-PGA (6.01 g, 40.0 mmol) in IPA (35 mL) and acetic acid (1.2 mL, 20 mmol) were stirred 4 days at 40 °C. The mixture was concentrated and the residue was dissolved in EtOAc (150 mL). The undissolved solids were filtered off and the filtrate was concentrated. Purification by flash chromatography (0-10% MeOH/ EtOAc) followed by crystallization afforded the enamine (5.47 g) as a white solid.

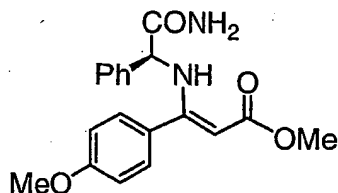
$[\alpha]_D^{25} = +18^\circ$ (c 0.33, DCM); mp = 111-130 °C.

¹H-NMR (DMSO-d₆, 400 MHz): δ 9.74 (d, *J* = 10 Hz, 1H), 7.58 (s, 1H), 7.33 (m, 3H), 7.22 (m, 3H), 7.16 (m, 2H), 7.08 (m, 3H), 6.9 (br, 1H), 6.3 (br, 1H), 4.79 (d, *J* = 10 Hz, 1H), 4.60 (s, 1H).
¹³C-NMR (DMSO-d₆, 100 MHz): δ 172.0, 171.5, 159.2, 140.9, 137.1, 129.2, 128.6, 128.5, 127.8, 127.6, 126.8, 92.8, 60.8.

5

EXAMPLE 10

Methyl (2Z)-3-([(1S)-2-amino-2-oxo-1-phenylethyl]amino)-3-(4-methoxyphenyl)acrylate



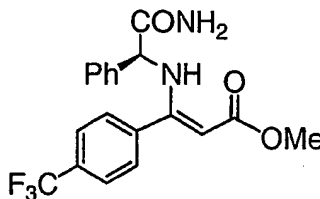
Methyl 3-(4-methoxyphenyl)-3-oxopropanoate (10.41 g, 50.0 mmol) and (*S*)-PGA (7.51 g, 50.0 mmol) in IPA (40 mL) and acetic acid (1.43 mL, 25 mmol) were stirred 5 days at 40 °C. The mixture was concentrated and EtOAc (50 mL) was added. The undissolved solids were filtered off and the filtrate was concentrated. The residue was purified by flash chromatography (30-100% EtOAc/ hexane) to afford the enamine as an amorphous solid, which was heated and stirred in MTBE (150 mL) to afford a slurry. The solids were filtered and dried to afford the enamine (5.96 g).

$[\alpha]_D^{25} = -31^\circ$ (c 0.56, DCM); mp = 93 °C (broad).

¹H-NMR (DMSO-d₆, 400 MHz): δ 9.41 (d, *J* = 8.9 Hz, 1H), 7.64 (s, 1H), 7.26 (m, 3H), 7.16 (m, 3H), 7.05 (d, *J* = 8.5 Hz, 2H), 6.89 (d, *J* = 8.6 Hz, 2H), 4.95 (d, *J* = 8.9 Hz, 1H), 4.50 (s, 1H), 3.75 (s, 3H), 3.60 (s, 3H).

¹³C-NMR (DMSO-d₆, 100 MHz): δ 171.2, 169.1, 162.6, 160.0, 140.1, 128.8, 128.3, 127.9, 127.4, 126.2, 113.7, 86.3, 60.1, 55.2, 49.9.

EXAMPLE 11
Methyl (2Z)-3-([(1S)-2-amino-2-oxo-1-phenylethyl]amino)-3-[4-(trifluoromethyl)phenyl]acrylate



Methyl 3-oxo-3-[4-(trifluoromethyl)phenyl]propanoate (5.18 g, 20.0 mmol) and (S)-PGA (3.00 g, 20.0 mmol) in IPA (25 mL) and acetic acid (1.2 mL, 20 mmol) were stirred 2 days at 40 °C. The mixture was cooled to RT, n-heptane (10 mL) was added to precipitate the unreacted PGA and filtered. The filtrate was concentrated and the residue was purified by flash chromatography (50-70% EtOAc/ hexane) to afford the enamine as an amorphous solid (6.2 g) which contained a small amount of the E-enamine isomer. The solids were crystallized from MTBE to afford the pure Z-enamine (4.18 g) as a white solid.

$[\alpha]_D^{25} = -79^\circ$ (c 0.57, DCM); mp = 91 °C (broad).

$^1\text{H-NMR}$ (DMSO- d_6 , 400 MHz): δ 9.43 (d, $J = 8.6$ Hz, 1 H), 7.70 (d, $J = 8$ Hz, 2 H), 7.60 (s, 1H), 7.30 (d, $J = 8$ Hz, 2H), 7.24 (~t, 4 H), 7.10 (d, $J = 6.5$ Hz, 2 H), 4.86 (d, $J = 8.6$ Hz, 1H), 4.57 (s, 1H), 3.62 (s, 3H).

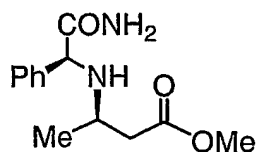
$^{13}\text{C-NMR}$ (DMSO- d_6 , 100 MHz): δ 170.8, 168.9, 160.9, 139.8, 139.6, 128.4, 128.3, 127.6, 126.2, 125.2, 87.3, 59.9, 50.1.

15 Examples of Hydrogenated PGA-Amines:

General Procedure: A pressure reaction vessel was charged with the PGA-enamine, PtO_2 and THF and the resulting slurry was cooled to 0 °C. The vessel was pressure-purged with nitrogen (3 x 40 psig) and hydrogen (2 x 40 psig) and then pressurized with hydrogen (90 psig). The reaction mixture was agitated for 30 min and then allowed to warm to room temperature over 15 min. After stirring for the prescribed amount of time, the vessel was carefully vented to atmospheric pressure and the resulting slurry was filtered through Solka floc. The filter cake was rinsed three times with THF and the filtrates were combined and concentrated. The residue was purified by flash chromatography (EtOAc/ hexane) and the product was crystallized (EtOAc/ hexane) when possible.

EXAMPLE 12

Methyl (3R)-3-[[[(1S)-2-amino-2-oxo-1-phenylethyl]amino]butanoate



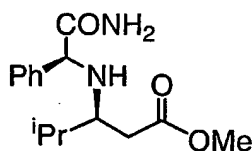
Flash chromatography (90% EtOAc/ hexane) afforded the ester.

¹H-NMR (DMSO-d₆, 400 MHz): δ 7.57 (bs, 1H), 7.45 (m, 2H), 7.39 (m, 2H), 7.33 (m, 1H), 7.20 (bs, 1H), 4.31 (bd, *J* = 4 Hz, 1H), 3.67 (s, 3H), 2.99 (m, 1H), 2.59 (dd, *J* = 15, 7 Hz, 2H), 2.38 (dd, *J* = 15, 7Hz, 2H), 1.11 (d, *J* = 6 Hz, 3H).

¹³C-NMR (DMSO-d₆, 100 MHz): δ 174.1, 172.2, 140.6, 128.1, 127.3, 127.2, 63.0, 51.2 47.9
5 41.3, 19.8.

EXAMPLE 13

Methyl (3*S*)-3-[[*(1S)*-2-amino-2-oxo-1-phenylethyl]amino]-4-methylpentanoate



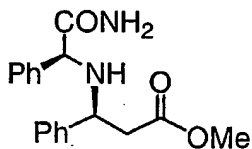
10 Flash chromatography (70-85% EtOAc/ hexane) afforded pure amine.

¹H-NMR (DMSO-d₆, 400 MHz): δ 7.42 (bs, 1H), 7.37 (m, 2H), 7.31 (m, 2H), 7.24 (m, 1H), 7.12 (bs, 1H), 4.20 (d, *J* = 6 Hz, 1H), 3.59 (s, 3H), 2.68 (m, 1H), 2.33 (m, 2H), 1.77 (m, 1H), 0.83 (d, *J* = 7 Hz, 3H), 0.76 (d, *J* = 7 Hz, 3H).

¹³C-NMR (DMSO-d₆, 100 MHz): δ 174.1, 172.8, 140.7, 128.0, 127.4, 127.1, 63.2, 57.4, 51.3,
15 29.7, 18.3, 17.4.

EXAMPLE 14

Methyl (3*R*)-3-[[*(1S)*-2-amino-2-oxo-1-phenylethyl]amino]-4-phenylbutanoate

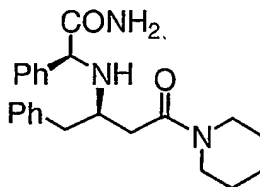


20 Flash chromatography (80-100% EtOAc/ hexane) afforded the amine. This was crystallized from EtOAc/ hexane.

¹H-NMR (DMSO-d₆, 400 MHz): δ 7.45 (s, 1H), 7.24 (m, 8H), 7.11 (m, 3H), 4.31 (d, *J* = 6.5 Hz, 1H), 3.55 (s, 3H), 2.97 (m, 1H), 2.80 (dd, *J* = 6.0, 13.3 Hz, 1 H), 2.58 (dd, *J* = 7.0 13.3 Hz, 1H), 2.37-2.46 (m, 2H), 2.29 (dd, *J* = 5.8, 15.2 Hz, 1H).

25 ¹³C-NMR (DMSO-d₆, 100 MHz): δ 173.8, 172.1, 140.2, 138.7, 129.2, 128.2, 128.0, 127.2, 127.2, 126.1, 62.8, 54.0, 51.4, 38.5.

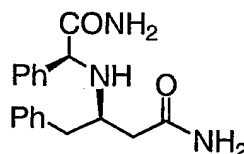
EXAMPLE 15

(2S)-2-[[[(1R)-1-benzyl-3-oxo-3-piperidin-1-ylpropyl]amino]-2-phenylacetamide

Flash chromatography (5% MeOH/ EtOAc) afforded the amine as an oil which was crystallized from 1:1 EtOAc/ hexane to afford the product as a white solid.

- 5 ¹H-NMR (DMSO-d₆, 400 MHz): δ 7.65 (s, 1H), 7.1-7.3 (m, 11H), 4.33 (d, *J* = 4.6 Hz, 1H), 3.39 (m, 2H), 3.20 (m, 2H), 2.98 (m, 1H), 2.81 (dd, *J* = 5.7, 13.2 Hz, 1H), 2.59 (dd, *J* = 6.9, 13.2 Hz, 1H), 2.3-2.5 (m, 2H), 2.22 (dd, *J* = 5.5, 15.2 Hz, 1H), 1.52 (m, 2H), 1.36 (m, 4H).
¹³C-NMR (DMSO-d₆, 100 MHz): δ 174.1, 169.1, 140.5, 139.2, 129.3, 128.2, 128.0, 127.3, 127.1, 126.0, 62.9, 54.2, 45.9, 41.8, 37.0, 26.0, 25.2, 23.9.

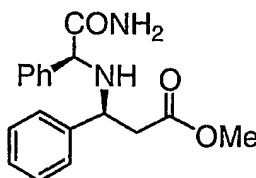
10

EXAMPLE 16(3R)-3-[[[(1S)-2-amino-2-oxo-1-phenylethyl]amino]-4-phenylbutanamide

- 15 Stirring the residue in 1:1 EtOAc/ hexane followed by filtration and drying afforded the amine as a white solid.

- ¹H-NMR (DMSO-d₆, 400 MHz): δ 7.64 (s, 1H), 7.46 (s, 1H), 7.25 (m, 8H), 7.11 (m, 3H), 6.83 (s, 1H), 4.36 (s, 1H), 2.97 (br, 1H), 2.80 (dd, *J* = 5.5, 13.2 Hz, 1H), 2.57 (dd, *J* = 7.0, 13.2 Hz, 1H), 2.28 (br, 1H), 2.10 (m, 2H).
¹³C-NMR (DMSO-d₆, 100 MHz): δ 174.1, 173.2, 140.5, 139.0, 129.3, 128.2, 128.0, 127.3, 127.2, 126.0, 62.8, 53.9, 39.4.

20

EXAMPLE 17Methyl (3S)-3-[[[(1S)-2-amino-2-oxo-1-phenylethyl]amino]-3-phenylpropanoate

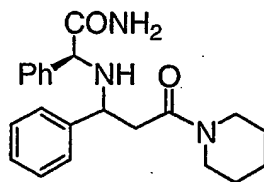
Flash chromatography (75% EtOAc/ hexane) afforded the pure amine.

$^1\text{H-NMR}$ (DMSO- d_6 , 400 MHz): δ 7.59 (d, J = 2.4 Hz, 1H), 7.2-7.3 (m, 11H), 3.98 (q, J = 7.7 Hz, 1H), 3.72 (d, J = 5.3 Hz, 1H), 3.56 (s, 3H), 3.14 (m, 1H), 2.86 (dd, J = 8.1, 15.4 Hz, 1H), 2.58 (dd, J = 6.4, 15.4 Hz, 1H).

5 $^{13}\text{C-NMR}$ (DMSO- d_6 , 100 MHz): δ 173.9, 171.6, 142.4, 140.1, 128.5, 128.2, 127.4, 127.4, 127.3, 127.2, 63.5, 57.6, 51.4, 42.3.

EXAMPLE 18

(2S)-2-[(3-oxo-1-phenyl-3-piperidin-1-ylpropyl)amino]-2-phenylacetamide



10

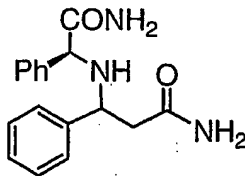
Flash chromatography (EtOAc/ hexane) afforded the amine as a white solid foam .

$^1\text{H-NMR}$ (DMSO- d_6 , 400 MHz): δ 7.99 (s, 1H), 7.36-7.19 (m, 11H), 4.03 (m, 1H), 3.70 (d, J = 5 Hz, 1H), 3.46 (m, 1H), 3.35 (m, 1H), 3.30 (m, 3H), 3.01 (bt, 1H), 2.81 (dd, J = 8.6, 15 Hz, 1H), 2.52 (dd, 4.9, 15 Hz, 1H), 1.52 (m, 2H), 1.39 (4 H).

15 $^{13}\text{C-NMR}$ (DMSO- d_6 , 100 MHz): δ 174.1, 168.6, 143.2, 140.5, 128.3, 128.1, 127.2, 127.2, 127.0, 63.6, 57.8, 46.0, 42.0, 40.7, 25.8, 25.2, 24.0.

EXAMPLE 19

3-[(1S)-2-amino-2-oxo-1-phenylethyl]amino}-3-phenylpropanamide

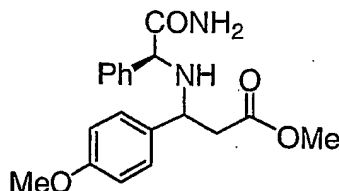


20

Suspension of the residue in EtOAc/ hexane and filtration afforded the product as a white solid.

$^1\text{H-NMR}$ (DMSO- d_6 , 400 MHz): δ 7.74 (s, 1H), 7.42 (s, 1H), 7.32 (m, 4H), 7.24 (m, 6H), 7.16 (s, 1H), 6.83 (s, 1H), 3.97 (bq, 1H), 3.74 (d, J = 5 Hz, 1H), 3.05 (bt, 1H), 2.52 (dd, J = 9.1, 15 Hz, 1H), 2.30 (dd, J = 5.1, 15 Hz, 1H).

25 $^{13}\text{C-NMR}$ (DMSO- d_6 , 100 MHz): δ 174.0, 172.5, 143.2, 140.4, 128.3, 128.1, 127.2, 127.1, 127.1, 63.3, 57.5, 43.7.

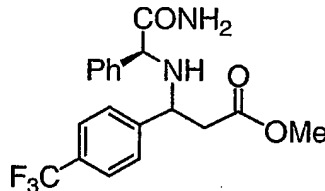
EXAMPLE 20Methyl 3-[[[(1S)-2-amino-2-oxo-1-phenylethyl]amino]-3-(4-methoxyphenyl)propanoate

Crystallization from EtOAc/ hexane afforded the amine as a white solid.

- 5 $^1\text{H-NMR}$ (DMSO- d_6 , 400 MHz): δ 7.55 (s, 1H), 7.23 (m, 8H), 6.89 (d, J = 8.6 Hz, 2H), 3.92 (br-q, 1H), 3.74 (s, 3H), 3.72 (1H), 3.55 (s, 3H), 3.06 (br-t, 1H), 2.84 (dd, J = 8, 15 Hz, 1H), 2.54 (dd, J = 6.5, 15 Hz, 1H).

$^{13}\text{C-NMR}$ (DMSO- d_6 , 100 MHz): δ 173.8, 171.5, 158.4, 140.1, 134.1, 128.2, 128.1, 127.2, 127.1, 113.7, 63.3, 56.8, 55.0, 51.2, 42.4.

10

EXAMPLE 21Methyl 3-[[[(1S)-2-amino-2-oxo-1-phenylethyl]amino]-3-[4-(trifluoromethyl)phenyl]propanoate

Flash chromatography (80% EtOAc/ hexane) afforded the amine.

- 15 $^1\text{H-NMR}$ (CD_3OD , 400 MHz): δ 7.66 (d, J = 8 Hz, 2H), 7.54 (d, J = 8 Hz, 2H), 7.27 (m, 5H), 4.25 (m, 1H), 3.88 (s, 1H), 3.68 (s, 3H), 2.89 (dd, J = 9.0, 16.0 Hz, 1H), 2.70 (dd, J = 5.6, 16.0 Hz, 1H).

$^{13}\text{C-NMR}$ (CD_3OD , 100 MHz): δ 178.0, 173.6, 148.0, 140.7, 129.8, 129.3, 129.2, 128.7, 126.8, 126.8, 65.6, 59.1, 52.4, 43.0

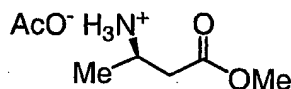
- 20 Examples of Deprotected Amines:

General Procedure: A pressure reaction vessel was charged with PGA-amine, acetic acid (2.5 molar equiv), water, THF, and 30% $\text{Pd}(\text{OH})_2$ (20 wt% based on PGA-amine) and warmed to 50 °C. The vessel was pressure-purged with nitrogen (3 x 40 psig) and pressurized with hydrogen (40 psig) and the reaction solution was stirred for 12 h. After cooling to room temperature, the
25 vessel was vented and the solution was filtered through Solka flocc. The filter cake was then

washed with THF. The filtrates were combined and the solvent was removed under reduced pressure to yield a mixture of the desired beta-amino acid derivative as the acetate salt and 2-phenylacetamide byproduct. The assay yield of the amine salt was determined by $^1\text{H-NMR}$ using added dichloromethane as an internal reference. The stereochemical assignment was made by measuring the optical rotation of the crude product as its HCl salt and comparing its sign with what is reported in the literature or of an authentic reference sample. The HCl salt was prepared by addition of HCl/ ether and evaporation to dryness.

EXAMPLE 22

10 Methyl (3R)-3-aminobutanoate acetate



$^1\text{H-NMR}$ (CD_3OD , 400 MHz): δ 3.73 (s, 3H), 3.65 (m, 1H), 2.70 (m, 2H), 1.95 (s, 3H), 1.34 (d, $J = 6.6$ Hz, 3H).

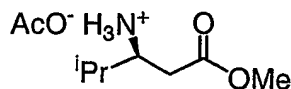
$^{13}\text{C-NMR}$ (CD_3OD , 100 MHz): δ 178.4, 172.5, 52.8, 45.6, 39.1, 23.0, 18.9.

15 Observed (HCl salt): $[\alpha]_D^{27} = \text{negative}$ (water) (Thus, stereochemistry is *R* as shown).

Literature for (*R*)-enantiomer (HCl salt): $[\alpha]_D^{28} = -37.0$ (c 0.73, water) [*J. Org. Chem.* **1992**, 57, 2396].

EXAMPLE 23

20 Methyl (3S)-3-amino-4-methylpentanoate acetate



$^1\text{H-NMR}$ (CD_3OD , 400 MHz): δ 3.74 (s, 3H), 3.38 (m, 1H), 2.76 (dd, $J = 4.0, 17.3$ Hz, 1H), 2.59 (dd, $J = 8.7, 17.3$ Hz, 1H), 1.96 (m, 1H), 1.93 (s, 3H), 1.01 (~t, 6H).

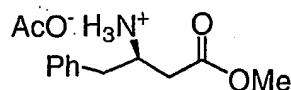
$^{13}\text{C-NMR}$ (CD_3OD , 100 MHz): δ 179.3, 173.4, 55.2, 53.1, 35.5, 32.3, 23.7, 19.0, 18.5.

25 Observed (HCl salt): $[\alpha]_D^{25} = \text{negative}$ (MeOH) (Thus, stereochemistry is *S* as drawn)

Literature for the (*R*)-enantiomer (HCl salt): $[\alpha]_D^{25} = +28.2$ (c 0.48, MeOH) [*Tetrahedron* **1995**, 51, 12337].

EXAMPLE 24

30 Methyl (3R)-3-amino-4-phenylbutanoate acetate



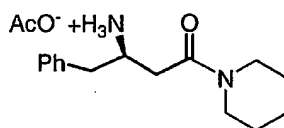
¹H-NMR (CD₃OD, 400 MHz): δ 7.3 (m, 5H), 3.71 (m, 1H), 3.69 (s, 3H), 2.64 (dd, *J* = 4.8, 17.0 Hz, 1H), 2.54 (dd, *J* = 7.8, 17.0 Hz, 1H), 1.94 (s, 3H).

¹³C-NMR (CD₃OD, 100 MHz): δ 172.9, 137.5, 130.6, 130.2, 128.6, 52.7, 51.0, 40.9, 37.9, 23.1.

- 5 Observed (HCl salt): [α]_D²⁵ = negative (MeOH) (Thus, stereochemistry is *R* as drawn).
Literature for the (*R*)-enantiomer (HCl salt): [α]_D = -9.2 (c 1.5, MeOH) [*Biosci. Biotech. Biochem.*, 1996, 60, 916].

EXAMPLE 25

- 10 [(1*R*)-1-benzyl-3-oxo-3-piperidin-1-ylpropyl]amine acetate



¹H-NMR (CD₃OD, 400 MHz): δ 7.3 (m, 5H), 3.78 (m, 1H), 3.52 (m, 2H), 3.32 (m, 2H), 3.08 (dd, *J* = 6.3, 13.7 Hz, 1H), 2.96 (dd, *J* = 8.5, 13.7 Hz, 1H), 2.64 (m, 2H), 1.96 (s, 3H), 1.63 (m, 2H), 1.51 (m, 4H).

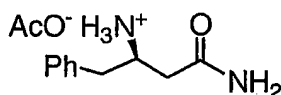
- 15 ¹³C-NMR (CD₃OD, 100 MHz): δ 177.9, 169.8, 137.3, 130.6, 130.2, 128.6, 51.5, 49.2, 47.6, 44.0, 39.6, 34.6, 27.4, 26.7, 25.4, 22.8.

Observed (HCl salt): [α]_D²⁰ = negative (MeOH) (Thus, stereochemistry is *R* as shown).

- An authentic sample of (*S*)-enantiomer (HCl salt) was prepared from Boc-L-β-homophenylalanine via EDC coupling with piperidine, followed by HCl deprotection: [α]_D²⁰ =
20 +37 (c 1.4, MeOH).

EXAMPLE 26

- (3*R*)-3-amino-4-phenylbutanamide acetate



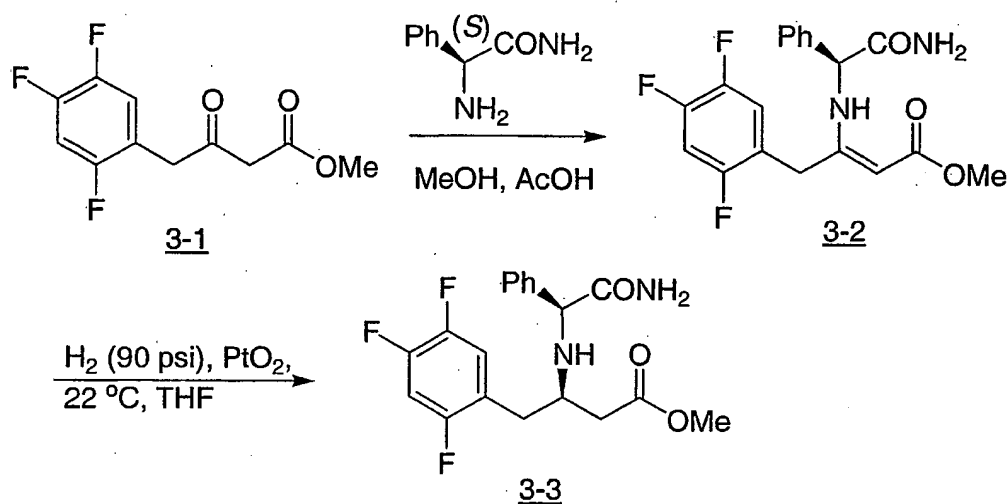
- 25 ¹H-NMR (CD₃OD, 400 MHz): δ 7.3 (m, 5H), 3.75 (m, 1H), 3.07 (dd, *J* = 6.1, 13.7 Hz, 1H), 2.90 (dd, *J* = 8.6, 13.7 Hz, 1H), 2.55 (m, 2H), 1.96 (s, 3H).

¹³C-NMR (CD₃OD, 100 MHz): δ 178.4, 175.2, 136.9, 130.6, 130.2, 128.6, 51.4, 39.7, 36.4, 23.0.

Observed (HCl salt): [α]_D²⁰ = negative (MeOH) (Thus, stereochemistry is *R* as shown).

An authentic sample of (*S*)-enantiomer (HCl salt) was prepared from Boc-L- β -homophenylalanine via EDC coupling with ammonia, followed by HCl deprotection: $[\alpha]_D^{20} = +11$ (c 1.0, MeOH).

5

EXAMPLE 27Step A: Preparation of Compound 3-2

A 2-liter 3-necked round-bottomed flask equipped with overhead stirrer, thermocouple, nitrogen inlet, and a condenser was charged with β -keto ester 3-1 (35 g), (*S*)-phenylglycine amide (21.35 g), methanol (700 mL) and acetic acid (44.78 mL). An exotherm of about 4 °C was observed during the acid addition. The mixture was heated to reflux. After 9 h, the mixture was removed from heating and allowed to cool to room temperature. When the temperature reached 40 °C, the product appeared as white solids. At 35 °C, the mixture turned into slurry. The mixture was stored at 4 °C. Concentration of the Z isomer in the supernatant was 7.08 mg/mL after 1 h and 3.19 mg/mL after overnight at 4 °C. Solids were filtered at 4 °C, washed with 140 mL MeOH and dried. The isolated solids were the Z isomer and have an *S*-configuration at the stereogenic center.

Step B: Preparation of Compound 3-3

The PGA enamine 3-2 (60 g), PtO_2 (6 g), acetic acid (45.4 mL), and THF (1.2 L) were charged into a flask. The mixture was hydrogenated for 15 h at 90 psi and room temperature. The catalysts were filtered through solka floc and rinsed with about 500 mL THF. Water (300 mL) was added and then the pH adjusted to about 7 with 50wt% NaOH over 15 min.

An exotherm of 9 °C was observed. The layers were then separated, and the organic layer was washed with water (150 mL). The organic layer was solvent switched at 45 °C to toluene (210 mL). The mixture was transferred to a round bottom flask equipped with mechanical stirrer. The product precipitated out as the solution cooled. The mixture was heated to 63 °C to redissolve the product, then cooled at 20 °C/h. It was seeded at 57 °C. After room temperature was reached, the mixture was cooled to 0 °C within 30 min and stored in a cold room overnight. The solids were filtered and washed with 60 mL toluene and dried under nitrogen sweep. Yield 50.25 g with 100 % diastereomeric excess

HPLC method:

10	Column	YMC Pro C18 (250 x 4.6 mm)
	Detector:	210 nm
	Mobile phase A	Acetonitrile
	Mobile phase B	10 mM K ₂ HPO ₄ pH 6.6 with H ₃ PO ₄
	Flow:	1.25 mL / min
15	Run Time:	15 min
	Gradient	Ramp from 50 % A to 60% A over 6 mins. Hold for 9 min.

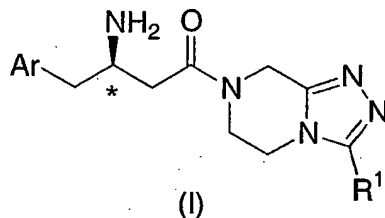
Typical Retention Times:

	Undesired isomer:	6.3 min
	Desired isomer:	6.5 min
20	PGA enamine <u>3-2</u> :	8.1 min

WHAT IS CLAIMED IS:

1. A process for preparing an enantiomerically enriched compound of structural formula I:

5

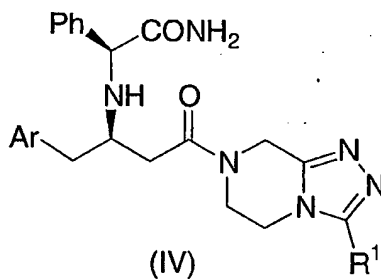


or a pharmaceutically acceptable salt thereof;

having the (*R*)-configuration at the stereogenic center marked with an *;

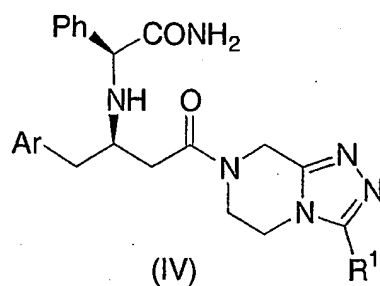
wherein

- 10 Ar is phenyl which is unsubstituted or substituted with one to five substituents independently selected from the group consisting of fluorine, trifluoromethyl, and trifluoromethoxy; and R¹ is hydrogen or C₁₋₄ alkyl unsubstituted or substituted with one to five fluorines; comprising the step of hydrogenolyzing a compound of structural formula IV:

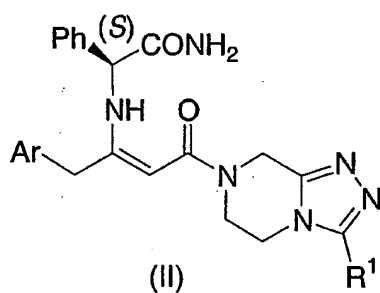


- 15 in the presence of a catalyst in a suitable organic solvent.

2. The process of Claim 1 additionally comprising the step of producing a compound of structural formula IV:



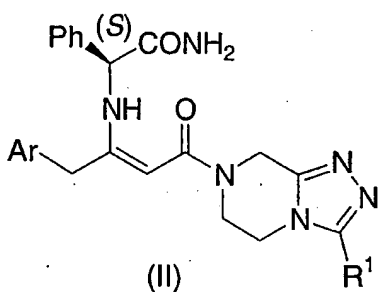
by hydrogenating a compound of structural formula II:



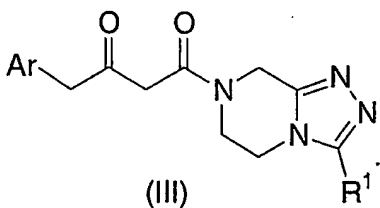
in the presence of a catalyst in a suitable organic solvent.

5

3. The process of Claim 2 additionally comprising the step of producing a compound of structural formula II:



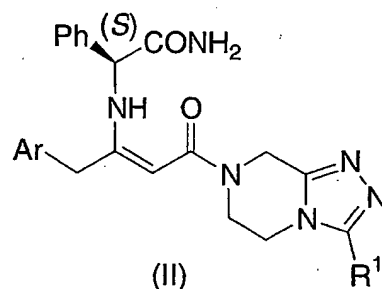
by treating a compound of structural formula III:



10

with (*S*)-phenylglycine amide in the presence of an acid in a suitable organic solvent.

4. The process of Claim 1 wherein said catalyst is a palladium catalyst.
5. The process of Claim 4 wherein said palladium catalyst is Pd(OH)₂/C.
6. The process of Claim 2 wherein said catalyst is platinum oxide.
7. The process of Claim 3 wherein said acid is acetic acid.
8. The process of Claim 1 wherein Ar is 2,5-difluorophenyl or 2,4,5-trifluorophenyl and R¹ is trifluoromethyl.
9. A compound of structural formula II:

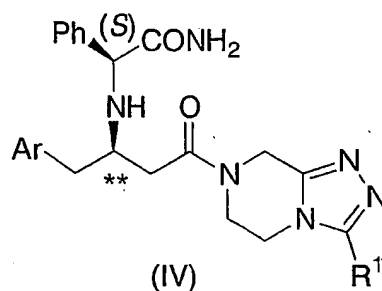


wherein

Ar is phenyl which is unsubstituted or substituted with one to five substituents independently selected from the group consisting of fluorine, trifluoromethyl, and trifluoromethoxy; and R¹ is hydrogen or C₁₋₄ alkyl unsubstituted or substituted with one to five fluorines.

10. The compound of Claim 9 wherein Ar is 2,5-difluorophenyl or 2,4,5-trifluorophenyl and R¹ is trifluoromethyl.

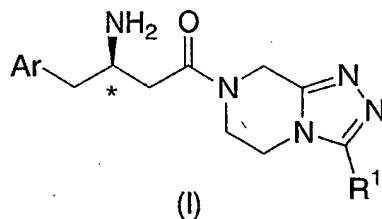
11. A compound of structural formula IV:



having the (*R*)-configuration at the stereogenic center marked with an **; wherein Ar is phenyl which is unsubstituted or substituted with one to five substituents independently selected from the group consisting of fluorine, trifluoromethyl, and trifluoromethoxy; and R¹ is hydrogen or C₁₋₄ alkyl unsubstituted or substituted with one to five fluorines.

12. The compound of Claim 11 wherein Ar is 2,5-difluorophenyl or 2,4,5-trifluorophenyl and R¹ is trifluoromethyl.

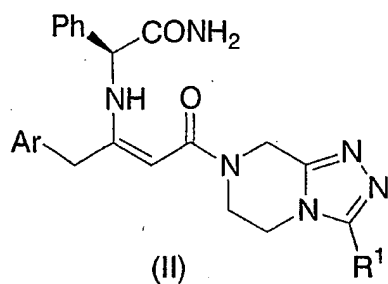
10 13. A process for the preparation of an enantiomerically enriched compound
of structural formula I:



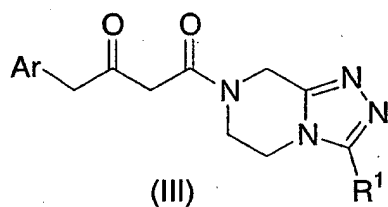
or a pharmaceutically acceptable salt thereof;
having the (*R*)-configuration at the stereogenic center marked with an *;

15 wherein
Ar is phenyl which is unsubstituted or substituted with one to five substituents independently
selected from the group consisting of fluorine, trifluoromethyl, and trifluoromethoxy; and
R¹ is hydrogen or C₁₋₄ alkyl unsubstituted or substituted with one to five fluorines;
comprising the steps of:

20 (a) producing a compound of structural formula II:

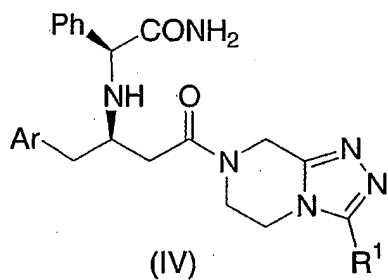


by treating a compound of structural formula III:

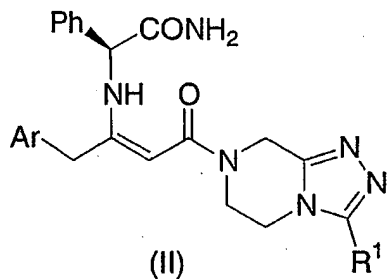


with (*S*)-phenylglycine amide in the presence of an acid in a suitable organic solvent;

5 (b) producing a compound of structural formula IV:

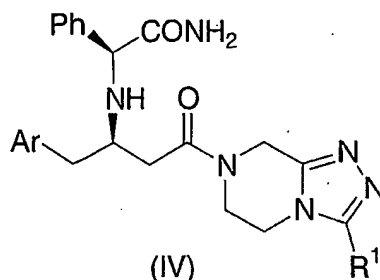


by hydrogenating a compound of structural formula II:



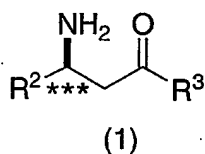
in the presence of a catalyst in a suitable organic solvent; and

10 (c) hydrogenolyzing a compound of structural formula IV:



in the presence of a catalyst in a suitable organic solvent to afford a compound of structural formula I.

- 5 14. A process for the preparation of an enantiomerically enriched beta-amino acid derivative of structural formula 1:



or an amine salt thereof;

having the indicated stereochemical configuration at the stereogenic center marked with an ***;

10 wherein

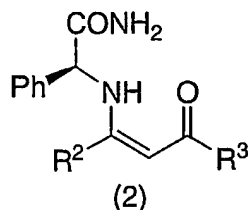
R² is C₁₋₈ alkyl, C₅₋₇ cycloalkyl, aryl, heteroaryl, aryl-C₁₋₂ alkyl, or

heteroaryl-C₁₋₂ alkyl, wherein aryl and heteroaryl are unsubstituted or substituted with one to three substituents independently selected from C₁₋₄ alkyl, halogen, C₁₋₄ alkoxy, and trifluoromethyl;

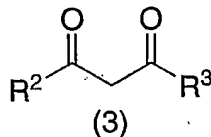
15 R³ is OR⁴, SR⁴, or NR⁴R⁵;

R⁴ and R⁵ are each independently hydrogen, C₁₋₈ alkyl, aryl, or aryl-C₁₋₂ alkyl; or R⁴ and R⁵ together with the nitrogen atom to which they are attached form a 4- to 7-membered heterocyclic ring system optionally containing an additional heteroatom selected from O, S, and NC₁₋₄ alkyl; comprising the steps of:

20 (a) producing a compound of structural formula 2:

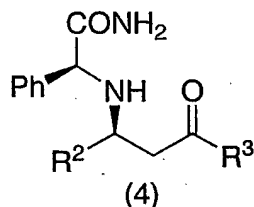


by treating a compound of structural formula 3:



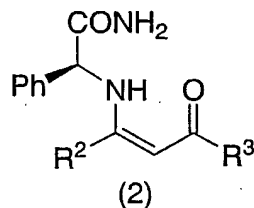
with (*S*)-phenylglycine amide in the presence of an acid in a suitable organic solvent;

(b) producing a compound of structural formula 4:



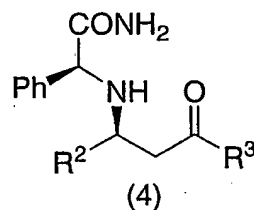
5

by hydrogenating a compound of structural formula 2:



in the presence of a catalyst in a suitable organic solvent; and

(c) hydrogenolyzing a compound of structural formula 4:



10

in the presence of a catalyst in a suitable organic solvent to afford a compound of structural formula 1 or a salt thereof.

15 15. The process of Claim 14 wherein R² is 2,4,5-trifluorobenzyl and R³ is C₁₋₄ alkoxy.

16. The process of Claim 15 wherein R³ is methoxy.